Attachment #8

The purpose of this study was to determine the potential of the PMN substance to cause acute dermal irritation in rabbits. The study was conducted in accordance with OECD Guideline 404. The PMN substance was applied to the skin of 3 rabbits using a gauze patch held in place with non-irritating tape for 4 hours. Skins were evaluated at 1, 24, 48 and 72 hours and at 7 days after dosing.

No edema was seen. Marginal erythema (GRADE 1) was seen through 72 hours. Full recovery was reported by day 7. The PMN substance was not classified as a dermal irritant in the assay. No systemic effects were noted.